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URETHANE ACRYLATE GEL COAT RESIN
AND METHOD OF MAKING

FIELD OF THE INVENTION

The present invention relates to improved
5 resins for use in gel coat compositions.

BACKGROUND OF THE INVENTION

Coated, molded articles, often fiber-
reinforced, typically are made by spreading a "gel
coat composition" over the surface of a mold having
10 a surface corresponding to the article in negative
relief. Consequently, the gel coat composition,
after cure, becomes the outermost layer of the
molded article that is exposed to the environment.
The gel coat composition is spread across the sur-
15 face of the mold by any one of a number of conven-
tional techniques, e.g., brushing, hand lay-up, or
spraying, and usually as a relatively thick layer,
e.g., 0.5 to 0.8 mm, to maximize its weather and
wear resistance, and if the molded article is fiber-
20 reinforced, to help mask the fiber reinforcement
pattern which can show through the gel coat due to
inherent resin shrinkage that occurs around the
fibers during cure.

A gel coat is a prepromoted resin, typ-
25 ically a polyester, and typically is pigmented.
After the gel coat is applied to the surface of the
mold, it is at least partially cured. A plastic,
optionally fiber-reinforced, then is applied to the
partially or fully cured gel coat by any one of a

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number of conventional techniques, and the resulting laminate structure cured. The cure can be promoted through the use of free radical polymerization processes.

5 In addition to imparting weather and wear resistance to the molded article, the gel coat also imparts cosmetic properties to the article. In many applications, particularly consumer applications such as automobile parts, shower stalls, bathtub
10 enclosures, and appliances, a high initial gloss and extended gloss retention are very desirable or necessary properties for the molded article. Present-day gel coats often exhibit a high gloss upon cure, but lose this gloss over time due to a variety of
15 environmental factors, such as sunlight, heat, cold, water, and corrosive chemicals, long before the end of the useful life of the article. Moreover, loss of gloss is often accompanied by the appearance of surface defects, such as cracks, coarseness, and
20 blisters, and these often are indicative of structural deterioration of the molded article itself.

 The use of unsaturated polyesters in admixture with unsaturated aromatic monomers, such as styrene, in gel coat compositions is well known in
25 the art. The unsaturated polyesters are prepared from the condensation of unsaturated acids or acid anhydrides with polyols. A common unsaturated acid is either maleic anhydride or fumaric acid. While not intending to be bound by theory, it is believed
30 that ester linkages formed from these ingredients exhibit poor hydrolysis resistance, and consequently

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the overall film performance of a coating film based on these polymers is relatively poor. Aromatic diacids, such as isophthalic acid, have been used to help improve the hydrolysis resistance of the film.

5 However, the presence of aromatic nuclei reduces the exterior durability of the coating film.

One high quality gel coat is an isophthalic acid/neopentyl glycol (IPA/NPG)-based unsaturated polyester diluted in styrene monomer. However, the
10 cured gel coats are rather soft materials of overall low chemical resistance and limited outdoor durability. The art has a need for a more durable gel coat because IPA/NPG gel coats can fade and chalk, even before the molded plastic article is sold.

15 Other gel coats presently in use include epoxy, urethane, and vinyl ester resins, particularly when greater flexibility and water resistance are desired. However, these materials also tend to fade and lose their gloss quickly, usually require
20 higher curing temperatures, and are much more difficult to use than the commonly available unsaturated polyester products. In addition, these resins are difficult to formulate into gel coat compositions having desirable physical properties, in-mold curing
25 times, and handling properties without the use of more than a nominal amount of styrene or similar volatile monomer as a reactive diluent. Moreover, because these diluents are the subject of numerous federal, state, and local regulations, manufacturers
30 of molded plastic articles prefer to use gel coat

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compositions that contain minimal styrene or similar volatile monomers.

In particular, vinyl esters formed from the reaction of an aromatic polyepoxide with an
5 unsaturated monocarboxylic acid have excellent hydrolysis resistance. However, the presence of aromatic nuclei and the necessary addition of high levels of unsaturated aromatic monomers to obtain a sprayable viscosity leads to unacceptable exterior
10 durability. Vinyl ester resins based on aliphatic polyepoxides exhibit poor hydrolysis resistance.

SUMMARY OF THE INVENTION

Present-day gel coat compositions fail to meet the requirements for weatherability, color
15 stability, and hydrolysis resistance for external applications, such as automotive applications. These requirements include no significant loss of gloss, change in color, or build-up of chalky oxidation products on the surface of the cured gel coat.

20 While not intending to be bound by theory, present-day gel coat compositions fail because of the chemistry used in preparing the base resin incorporated into the gel coat composition. Typically, the chemistry is based on an unsaturated polyester, or a hybrid chemistry based on polyesters and
25 acrylates. The present invention is directed to a new resin for use in a gel coat composition that overcomes problems and disadvantages associated with prior base resins used in gel coat compositions.

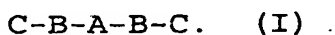
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The present invention, therefore, is directed to a urethane acrylate resin that exhibits substantially improved performance over present-day base resins used in gel coat compositions. Gel coat compositions containing a present urethane acrylate resin retain a high gloss and consistent color over extended time periods.

The above-described deficiencies in prior gel coat compositions have been overcome by incorporating a urethane acrylate resin of the present invention into a gel coat composition. The improved gel coat compositions provide cured gel coats having excellent weathering and hydrolytic stability.

In particular, the present invention is directed to a urethane acrylate gel coat resin. More particularly, the present invention is directed to a urethane acrylate gel coat resin that is a reaction product of (a) an oligoester of weight average molecular weight (M_w) about 200 to about 4000, (b) a diisocyanate, and (c) a hydroxyalkyl (meth)acrylate.

Accordingly, one important aspect of the present invention is to provide a urethane acrylate gel coat resin containing a reaction product of component A (oligoester), component B (diisocyanate), and component C (hydroxyalkyl (meth)acrylate), and having an idealized structure (I):



Notably, the reaction product of components A, B, and C also contains other species in addition to

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idealized structure (I) and this invention is not limited to idealized structure (I).

Another aspect of the present invention is to provide a urethane acrylate gel coat resin for
5 incorporation into a gel coat composition. The gel coat composition provides a cured gel coat having improved weatherability, including gloss retention and color stability.

Still another aspect of the present inven-
10 tion is to provide a urethane acrylate gel coat resin having terminal acrylate groups. The terminal acrylate groups can be polymerized, for example, using free radical polymerization techniques to provide a cured gel coat.

15 Another aspect of the present invention is to provide a urethane acrylate gel coat resin suitable for use in gel coat compositions, wherein the resin is the reaction product of (a) a hydroxy terminated oligoester having M_w of about 200 to about
20 4000, (b) a diisocyanate (preferably predominantly an aliphatic diisocyanate), and (c) a hydroxyalkyl (meth)acrylate, wherein a reaction mixture of (a), (b), and (c) has a molar ratio of about 0.75 to about 1.25 moles of (a), about 1.5 to about 2.5
25 moles of (b), and about 1.5 to about 2.5 moles of (c). Preferred mole ratios of (a), (b), and (c) are about 0.9 to about 1.1 moles (a), about 1.7 to about 2.5 moles (b), and about 1.7 to about 2.2 moles (c), and especially about 0.95 to about 1.05 moles (a),
30 about 1.7 to about 2 moles (b), and about 1.7 to about 2 moles (c).

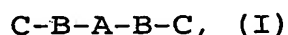
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These and other aspects and advantages of the present invention will become apparent from the following detailed description of the preferred embodiments.

5 **DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS**

The present invention is directed to a urethane acrylate gel coat resin useful as a base resin in gel coat compositions. After curing, a gel coat composition containing a resin of the present invention possesses not only very desirable gloss and gloss retention properties, but also exhibits excellent outdoor durability, hardness, toughness, and good handling properties during the molding process.

15 A urethane-acrylate gel coat resin of the present invention has an idealized structure (I)



wherein (I) is the reaction product of an oligoester having M_w of about 200 to about 4,000 (A), a diisocyanate (B), and a hydroxyalkyl (meth)acrylate (C). A urethane acrylate gel coat resin of the present invention is a reaction product of A, B, and C, thus other reactions species generally are present in addition to a resin of idealized structure (I).

25 In accordance with an important feature of the present invention, a present urethane acrylate gel coat resin contains an oligoester of M_w about 200 to about 4000 that is reacted with a diisocyanate, and the resulting urethane product is end-capped

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with a hydroxyalkyl (meth)acrylate. The urethane acrylate resin therefore contains terminal vinyl groups available for free radical polymerization, typically using a peroxide catalyst.

5 The individual ingredients used in the manufacture of a present urethane acrylate gel coat resin are described in more detail below.

(a) Oligoester

10 The oligoester component (A) of a present urethane acrylate gel coat resin preferably has a weight average molecular weight of about 200 to about 4000 and preferably is prepared from one or more saturated polyol and one or more saturated or unsaturated polycarboxylic acid or dicarboxylic acid
15 anhydride. As used herein, the terms "polyol" and "polycarboxylic" are defined as compounds that contain two or more, and typically two to four, hydroxy (OH) groups, or two or more, typically two or three, carboxyl (COOH) groups, respectively. Preferably,
20 the oligoester is hydroxy terminated to provide reactive moieties for a subsequent reaction with a diisocyanate.

 The polyesters typically are prepared from an aliphatic dicarboxylic acid or aliphatic dicar-
25 boxylic acid anhydride, and an aliphatic polyol. These ingredients are interacted preferably to provide a polyester having M_w of about 200 to about 4000, more preferably about 400 to about 3500, and most preferably about 500 to about 3000. Accord-

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ingly, the polyesters are low molecular weight oligoesters.

The oligoester typically is prepared, for example, by condensing an aliphatic dicarboxylic acid or aliphatic dicarboxylic acid anhydride with a polyol, preferably a diol. The polyol and dicarboxylic acid or acid anhydride, in correct proportions, are interacted under standard esterification procedures to provide an oligoester having the necessary M_w , molecular weight distribution, branching, and hydroxy-terminated functionality for use in a present urethane acrylate gel coat resin. In particular, the relative amounts of dicarboxylic acid and polyol are selected such that a sufficient excess molar amount of the polyol is present in order to provide a hydroxy terminated oligoester.

Nonlimiting examples of diols used to prepare the oligoesters include ethylene glycol, diethylene glycol, trimethylene glycol, propylene glycol, dipropylene glycol, hexylene glycol, 1,3-butylene glycol, 1,4-butylene glycol, neopentyl glycol, cyclohexanedimethanol, pinacol, pentanediol, 2,2-dimethyl-1,3-propanediol, isopropylidene bis(p-phenyleneoxypropanol-2), a polyethylene or polypropylene glycol having a weight average molecular weight of about 500 or less, and mixtures thereof. A small amount of a triol or polyol, e.g., up to 5 mole %, more preferably 0 to 3 mole % of a triol or polyol, can be used to provide a partially branched, as opposed to linear, oligoester. Nonlimiting exam-

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ples of a triol include glycerol and trimethylolpropane.

Exemplary dicarboxylic acids, and anhydrides thereof, used to prepare a hydroxy-terminated oligoester include aliphatic dicarboxylic acids, such as, but not limited to, adipic acid, malonic acid, cyclohexanedicarboxylic acid, sebacic acid, azelaic acid, succinic acid, glutaric acid, and mixtures thereof. Substituted aliphatic dicarboxylic acids, such as halogen or alkyl-substituted dicarboxylic acids, also are useful.

Additional suitable dicarboxylic acids, and anhydrides thereof, include maleic, dihydroxymaleic, diglycollic, oxalacetic, oxalic, pimelic, suberic, chlorosuccinic, mesoxalic, acetone dicarboxylic, dimethyl malonic, 1,2-cyclopropanedicarboxylic, cyclobutane-1,1-dicarboxylic, cyclobutane-1,2-dicarboxylic, cyclobutane-1,3-dicarboxylic, cyclopentane-1,1-dicarboxylic, cyclopentane-1,2-dicarboxylic, 2,5-dimethylcyclopentane-1,1-dicarboxylic, alpha,alpha'-di-sec-butyl-glutaric, beta-methyl-adipic, isopropyl-succinic, and 1,1-dimethyl-succinic acids.

Additional suitable diols, triols, polyols, dicarboxylic acids and anhydrides, and polycarboxylic acids are disclosed in U.S. Patent No. 5,777,053, incorporated herein by reference.

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(b) Diisocyanate

The diisocyanate component (B) of a present urethane acrylate gel coat resin is an aliphatic diisocyanate. The diisocyanate component optionally
5 can contain up to about 20%, and preferably up to about 10%, by total weight of the diisocyanate, of an aromatic diisocyanate. The identity of the aliphatic diisocyanate is not limited, and any commercially available commercial or synthetic diiso-
10 cyanate can be used in the manufacture of a urethane acrylate gel coat resin of the present invention.

Nonlimiting examples of aliphatic diisocyanates include 1,6-hexamethylene diisocyanate, isophorone diisocyanate, 1,4-cyclohexane diiso-
15 cyanate, 2,4'-dicyclohexylmethane diisocyanate, 4,4'-dicyclohexylmethane diisocyanate, 1,3-bis(isocyanatomethyl)cyclohexane, 1,4-bis(isocyanatomethyl)cyclohexane, tetramethylxylylene diisocyanate, 1,11-diisocyanatoundecane, 1,12-diiso-
20 cyanatododecane, 2,2,4-trimethyl-1,6-diisocyanatohexane, 2,4,4-trimethyl-1,6-diisocyanatohexane, 1,2-bis(isocyanatomethyl)cyclobutane, hexahydro-2,4-diisocyanatotoluene, hexhydro-2,6-diisocyanatotoluene, 1-isocyanato-2-isocyanatomethyl cyclopentane, 1-isocyanato-3-isocyanatomethyl-3,5,5-trimethyl cyclohexane, 1-isocyanato-4-isocyanatomethyl-1-methyl cyclohexane, 1-isocyanato-3-isocyanatomethyl-1-methyl cyclohexane, and mixtures thereof. A preferred aliphatic diisocyanate is isophorone diiso-
30 cyanate.

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Nonlimiting examples of optional aromatic diisocyanates includes toluene 2,4-diisocyanate, toluene 2,6-diisocyanate, 4,4'-methylene diphenyl diisocyanate, 2,4'-methylene diphenyl diisocyanate, 5 polymeric methylene diphenyl diisocyanate, p-phenylene diisocyanate, naphthalene-1,5-diisocyanate, and mixtures thereof.

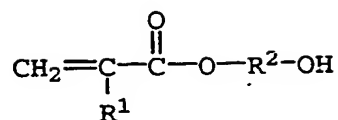
Additional aliphatic and aromatic diisocyanates are disclosed in U.S. Patent No. 5,777,053, 10 incorporated herein by reference.

(c) Hydroxyalkyl (meth)acrylate)

The hydroxyalkyl (meth)acrylate component (C) of a present urethane acrylate gel coat resin is preferably a hydroxyalkyl ester of an α,β -unsaturated acid, or anhydride thereof. Suitable α,β -unsaturated acids include a monocarboxylic acid such as, but not limited to, acrylic acid, methacrylic acid, ethacrylic acid, α -chloroacrylic acid, α -cyanoacrylic acid, β -methylacrylic acid (crotonic acid), α -phenylacrylic acid, β -acryloxypropionic acid, cinnamic acid, p-chlorocinnamic acid, β -stearylacrylic acid, and mixtures thereof. As used throughout this specification, the term "(meth)acrylate" is an abbreviation for acrylate and/or 25 methacrylate.

A preferred acrylate monomer containing a hydroxy group is a hydroxyalkyl (meth)acrylate having the following structure:

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wherein R¹ is hydrogen or methyl, and R² is a C₁ to C₆ alkylene group or an arylene group. For example, R² can be, but is not limited to (-CH₂-)_n, wherein n is
 5 1 to 6,



any other structural isomer of an alkylene group containing three to six carbon atoms, or can be a cyclic C₃-C₆ alkylene group. R² also can be an arylene group like phenylene (i.e., C₆H₄) or naphthylene (i.e., C₁₀H₆). R² optionally can be substituted with relatively nonreactive substituents, like C₁-C₆ alkyl, halo (i.e., Cl, Br, F, and I), phenyl, alkoxy, and aryloxy (i.e., an OR² substituent).
 10

Specific examples of monomers containing a hydroxy group are the hydroxy(C₁-C₆)alkyl (meth)acrylates, e.g., 2-hydroxyethyl methacrylate, 2-hydroxyethyl acrylate, 2-hydroxypropyl methacrylate, and 3-hydroxypropyl methacrylate.
 15

20

The relative amounts of (a), (b), and (c) used in the manufacture of a urethane acrylate gel coat resin of the present invention are sufficient to provide a reaction product having an idealized structure (I). Accordingly, component (a) is used
 25 in a molar amount of about 0.75 to about 1.25, and

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preferably about 0.9 to 1.1 moles; component (b) is used in an amount of 1.5 to about 2.5, and preferably about 1.7 to about 2.2 moles; and component (c) is used in an amount of about 1.5 to about 2.5, and preferably about 1.7 to about 2.2 moles. To achieve the full advantage of the present invention, the mole ratio of (a):(b):(c) is 1:1.7-2:1.75-2.

A urethane acrylate gel coat resin of the present invention is manufactured by first preparing the oligoester. The oligoester is prepared from a polyol, predominantly or completely a diol, and a polycarboxylic acid, predominantly or completely a dicarboxylic acid or anhydride thereof, using standard esterifying condensation conditions. The amounts and relative amounts of polyol and polycarboxylic acid are selected, and reaction conditions are used, such that the oligoester preferably has an M_w of about 200 to about 4000 and is hydroxy terminated. The oligoester can be saturated or unsaturated.

The oligoester then is blended with the hydroxyalkyl (meth)acrylate, followed by addition of the diisocyanate. The resulting reaction leads to a mixture of products, including a species having the idealized structure (I). Structure (I) has terminal acrylate moieties available for polymerization using standard free radical techniques, e.g., using initiators such as peroxides or peroxy esters.

To demonstrate the usefulness of a urethane acrylate gel coat resin of the present invention, the following examples were prepared. These

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resins can be incorporated into a gel coat composition, which, after curing, exhibits excellent weatherability and color stability.

The following abbreviations are used in the Examples:

NPG	neopentyl glycol
MA	maleic anhydride
DBTDL	Dibutyl tin dilaurate
HEA	2-hydroxyethyl acrylate
IPDI	isophorone diisocyanate
MMA	methyl methacrylate
THQ	toluhydroquinone
TMP	trimethylolpropane
HALS	hindered amine light stabilizer
BYK-A-555	silicone defoamer, commercially available from BYK-Chemie USA, Inc.
AEROSIL 200	fumed silica, commercially available from Degussa Corporation
SARTOMER SR-9021	
SARTOMER SR-206	
DMAA	dimethyl acetoacetamide
TINUVAN 928	2-(2H-benzotriazol-2-yl)-6-(1-methyl-1-phenylethyl-4-(1,1,3,3-tetramethylbutyl)-phenol, commercially available from Ciba Specialty Chemicals Corporation
TINUVAN 123	bis-(1-octyloxy-2,2,6-tetramethyl-4-piperidyl) sebacate, a commercially available HALS from Ciba Specialty Chemicals Corporation

EXAMPLE 1

NPG (101.64 wt. parts), MA (60.59 wt. parts), and DBTDL (0.42 wt. parts) were added into a flask equipped with a packed column and agitator. The resulting mixture was heated to a maximum of 440°F and reacted to an acid number of about 5-10 under a nitrogen atmosphere by removing water (11.14 wt. parts). To the resulting oligoester (151.65 wt.

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parts) was added 2,6-di-t-butyl-p-cresol (0.65 wt. parts) and HEA (75.71 wt. parts) at 200°F. IPDI (114.28 wt. parts) was added to the resulting mixture via an addition funnel to maintain the exothermic reaction temperature below 200°F. The reaction was maintained at 200°F for one hour followed by the addition of MMA (107.69 wt. parts) as a solvent and THQ (0.03 wt. parts) as an inhibitor. The resulting product was 80%, by weight, urethane acrylate gel coat resin in 20%, by weight, MMA solvent.

EXAMPLE 2

The urethane acrylate gel coat resin of this example contains a saturated oligoester. As in Example 1, the oligoester is reacted with IPDI and HEA to produce a urethane polyester copolymer having acrylic unsaturation at the terminal positions. The resin of Example 2 is prepared in a manner essentially identical to Example 1.

Ingredient	Moles	Wt. Parts
1. 1,6-Hexanediol	2.69	24.76
2. TMP	0.07	0.68
3. Adipic acid	2	22.66

Ingredients 1-3 were reacted under esterifying conditions to remove 5.78 wt. parts of water, and provide an oligoester (40.78 wt. parts) of equivalent weight 239.1. The following ingredients were added to the oligoester, and reacted to form a

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urethane acrylate gel coat resin of the present invention.

Ingredient	Moles	Wt. Parts
4. DBTDL		0.08
5. 2,6-di-t-butyl-p-cresol		0.13
6. HEA	2.11	13.92
7. IPDI	4	25.30
8. THQ		0.006
9. MMA		19.79

5

EXAMPLE 3

1,6-Hexanediol (94.8 wt. parts) and TMP (2.6 wt. parts) were added into a flask equipped with an agitator, and the mixture was melted. Next, adipic acid (86.8 wt. parts) was added, and the resulting mixture was heated to 440°F, under a nitrogen atmosphere. An esterification reaction was performed, at a maximum temperature of 460°F, until the acid number was less than 10, preferably less than 7. Water (21.1 wt. parts) was removed during the reaction. The resulting oligoester was cooled to 140°F using a one part air sparge and 2 part nitrogen blanket. Next, DBTDL (0.31 wt. parts), 2,6-di-t-butyl-p-cresol (0.53 wt. parts), HEA (55.7 wt. parts), and IPDI (101.2 wt. parts) were added to the oligoester. The IPDI was added at a rate such that the exothermic reaction was maintained below 200°F (e.g., over about 30-60 minutes). The reaction was continued for 2 to 3 hours, periodically testing for free isocyanate groups (% NCO). A % NCO of less than 0.3 is preferred. At the completion of

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the reaction, THQ (0.03 wt. parts) and MMA (79.2 wt. parts) were added slowly to the urethane acrylate gel coat resin at a temperature below 190°F. The resulting mixture was stirred at 140°F for at least one hour. The resulting product contained 80% urethane acrylate gel coat resin and 20% MMA solvent.

The urethane acrylate gel coat resins of the present invention can be used in gel coat compositions. A resin of the present invention is the base resin of the gel coat composition, and can be formulated with other standard gel coat composition ingredients. The urethane acrylate gel coat resin can be cured by polymerization of the terminal acrylate groups using standard free radical techniques.

In particular, gel coat compositions can be formulated using the resins of this invention in the usual method. Gel coat compositions include pigments, extenders, promoters, catalysts, stabilizers, and the like as practiced in the art. Such gel compositions typically comprise about 25 to about 50 weight percent urethane acrylate gel coat resin, and about 10 to about 50 weight percent styrene or other vinyl monomer, said percentages being based on combined weights of resin and vinyl monomer. Other gel coat composition-ingredients include acrylic diluents (e.g., MMA), additives (e.g., silica, cobalt salts, silicone release agent, hydroxyalkyl (meth)acrylates, dimethyl acetoaceto-

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(e.g., methyl ethyl ketone peroxide), UV stabilizers, thixotropes, and other resins (e.g., an isophthatic-NPG-maleic unsaturated polyester).

The preparation of a gel coat composition, and curing of a gel coat composition to provide a gel coat for an article of manufacture are generally disclosed in WO 94/07674 and U.S. Patent No. 4,742,121, incorporated herein by reference. Gel coat compositions incorporating a urethane acrylate gel coat resin of the present invention are disclosed in Kia et al. U.S. provisional application entitled "Gel Coat Composition," filed on August 9, 2002 (GM Ref. No. GP-301493, HD&P Ref. No. 8540R-000005), incorporated herein by reference.

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Example 4 General Dark Color Gel Coat Formula	
	wt. %
Urethane acrylate gel coat resin (80% in MMA)	38-50
Styrene	0-5
Air release agent	.1-1
Thixotrope	.5-3
Reactive monomer	20-35
Cobalt	.1-.5
Cobalt promoter	.2-.7
UV inhibitor	.2-.5
HALS	.2-1
Glycol synergist	.1-1.5
Pigment paste	10-25

Fillers (e.g., mica, aluminum trihydrate, barium sulfate, and the like) are optional ingredients present at 0-15 wt.%. Blocked isocyanates are also optional ingredients present at 0-20 wt.%. 20

Examples of reactive monomers include, but are not limited to, methyl methacrylate (10-20

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wt.%), ethylene glycol dimethacrylate, e.g., SARTOMER SR-206 (1-10 wt.%), highly propoxylated glyceryl triacrylate, e.g., SARTOMER SR-9021 (0-10 wt.%), and mixtures thereof.

- 5 The pigment paste contains a pigment in an unsaturated polyester carrier resin. The paste also contains wetting agents, dispersing agents, and inhibitors, in minor amounts. Saturated polyesters also can be used as the carrier resin. The carrier
- 10 resin also can be different from a polyester, e.g., a urethane diacrylate, an acrylic silicone, or similar resin. The pigment paste is prepared by adding the pigment and other ingredients to the carrier resin, then mixing in a grinding machine.

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Example 5 Blue Gelcoat Composition	
Ingredient	Weight (kg)
Urethane acrylate gel coat resin of Example 2	42
Styrene monomer	4
BYK-A 555	1
AEROSIL 200	2
Grind to 6 on Hegmann gauge	
SARTOMER SR 9021	10
SARTOMER SR-206	1
Methyl methacrylate	19
Cobalt octoate (12%) in mineral spirits and dipropylene glycol monomethyl ether	0.5
DMAA	0.1 (gram)
TINUVIN 928	.5
TINUVIN 123	1
2-hydroxyethyl methacrylate	1
Blend 10 minutes	
Blue tinter	17
White tinter	1

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Example 6 White Gelcoat Composition	
Ingredient	Weight (kg)
Urethane acrylate gel coat resin of Example 2	24.6149
Styrene monomer	4
BYK-A 555	1
AEROSIL 200	.5
Grind to 6 on Hegmann Gauge	
SARTOMER SR-9021	7
SARTOMER SR-206	1
Methyl methacrylate	17.175
Cobalt octoate (12%) in mineral spirits and dipropylene glycol monomethyl ether	.2
DMAA	.1 gram
TINUVIN 928	.5
TINUVIN 123	1
2-Hydroxyethyl methacrylate	1
Blend 10 minutes	
Blue tinter	.01
White tinter	42

A gel coat composition comprising a urethane acrylate gel coat resin of the present invention, after curing, exhibits excellent weatherability and color stability. The urethane acrylate gel coat resin also is readily formulated into gel coat composition. Furthermore, incorporation of a urethane acrylate gel coat resin of the present invention into a gel coat composition permits a significant reduction in the amount of other resins, such as unsaturated polyesters, that typically are included in the gel coat composition. The elimination or reduction of unsaturated polyesters helps improve the weatherability and color stability of cured gel coats.

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The above-described advantages of a present urethane acrylate gel coat resin provide an improved gel coat composition useful for application as an exterior of a molded article, for example, an automobile part, an appliance, a bathtub, a shower stall, and similar reinforced plastic articles of manufacture. A urethane acrylate gel coat resin of the present invention can be used in a variety of gel coat compositions, and, therefore, has a wide range of applications. The enhanced performance characteristics of a gel coat composition comprising a present urethane acrylate gel coat resin is achieved by a novel combination of ingredients utilized to manufacture the urethane acrylate gel coat resin.

Obviously, many modifications and variations of the invention as hereinbefore set forth can be made without departing from the spirit and scope thereof, and, therefore, only such limitations should be imposed as are indicated by the appended claims.